## CLAIMS:

- 1. A method for synthesizing a peptide dimer, comprising:
- (a) providing a linking moiety L<sub>K</sub> having first and second functional groups capable
  of serving as initiation sites for peptide synthesis, and a third functional group attachable to a solid support;
  - (b) binding the linking moiety  $L_K$  to a solid support through the third functional group;
  - (c) synthesizing a first peptide segment at the first functional group and a second peptide segment at the second functional group, wherein each of said first and second peptide segments contain two cysteine residues positioned to allow intramolecular cyclization through a disulfide bond;
  - (d) oxidizing the compound provided in step (c) in a manner effective to promote formation of disulfide bonds between cysteine residues in the same peptide segment while minimizing formation of disulfide bonds between cysteine residues in different peptide segments.
  - 2. The method of claim 1, wherein step (d) comprises treatment with an oxidizing composition containing an oxidizing reagent of a type and in an amount effective to minimize reaction products in which a cysteine residue of the first peptide segment binds to a cysteine residue of the second peptide segment.
    - 3. The method of claim 2, wherein the oxidizing reagent is dimethyl sulfoxide.
- 4. The method of claim 3, wherein the oxidizing composition comprises approximately 15% to 100% (v/v) dimethyl sulfoxide.
  - 5. The method of claim 4, wherein the oxidizing composition comprises approximately 50% to 100% (v/v) dimethyl sulfoxide.
  - 6. The method of claim 5, wherein the oxidizing composition comprises approximately 50% to 100% (v/v) dimethyl sulfoxide.

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- 7. The method of claim 6, wherein the oxidizing composition comprises approximately 80% to 100% (v/v) dimethyl sulfoxide.
- 8. The method of claim 7, wherein the oxidizing composition comprises approximately 100% (v/v) dimethyl sulfoxide.

## 9. The method of claim 1, wherein:

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the first peptide segment is approximately 10 to 40 amino acid residues in length, binds to the erythropoietin receptor, and contains a sequence of amino acids  $X_3X_4X_5GPX_6TX_7X_8X_9$  (SEQ ID NO: 1) wherein each amino acid is indicated by standard one-letter abbreviation,  $X_3$  is C or homocysteine (Hoc),  $X_4$  is R, H, L or W,  $X_5$  is M, F, I or nor-leucine (J),  $X_6$  is selected from any one of the 20 genetically coded L-amino acids and J,  $X_7$  is W, 1-naphthylalanine (B) or 2-naphthylalanine (U),  $X_8$  is D, E, I, L or V, and  $X_9$  is C or Hoc; and

the second peptide segment is approximately 10 to 40 amino acid residues in length, binds to the erythropoietin receptor, and contains a sequence of amino acids  $X'_3X'_4X'_5GPX'_6TX'_7X'_8X'_9$  (SEQ ID NO: 2) wherein each amino acid is indicated by standard one-letter abbreviation,  $X'_3$  is C or Hoc,  $X'_4$  is R, H, L or W,  $X'_5$  is M, F, I or J,  $X'_6$  is selected from any one of the 20 genetically coded L-amino acids and J,  $X'_7$  is W, B or U,  $X_8'$  is D, E, I, L or V, and  $X'_9$  is C or Hoc.